RESPONSE TO OFFICE ACTION

ATTY. DOCKET : RM.CH5

APPLICANT : Symington, et al.
US SERIAL NO. : 10/539,923
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Art Unit : 1612 Conf. No. : 90068399

Annexure 1 - Claims Rewritten to Show Amendments

Please amend the claims to read as follows:

1. (Currently Amended) A formulation for a pharmacologically inert coating to serve as a temporary mechanical barrier on top of a temporary coating of a pharmacologically active substance be applied to a surface of a tooth of a patient, the formulation comprising, in combination:

a pharmacologically-active substance applied directly onto a surface of the tooth of the patient; and

a coating applied over the pharmacologically-active substance, said coating being formed of;

an aqueous dispersion of a polymethylmethacrylate; and a plasticizer.

- 2. (Currently Amended) The formulation of claim 1 wherein the <u>aqueous dispersion</u>
 of the polymethylmethacrylate is comprises an ammonio methacrylate copolymer, type B
 USP/NF.
- 3. (Currently Amended) The formulation of claim 2 wherein the ammonio methacrylate copolymer is EUDRAGIT RS 30 D brand polymethylmethacrylate.
- 4. (Original) The formulation of claim 1 wherein the plasticizer is a pharmaceutical grade plasticizer selected from the group consisting of triethyl citrate, dibutyl sebacate, dibutyl phthalate, and diethyl phthalate.
 - 5. (Original) The formulation of claim 4 wherein the plasticizer is triethyl citrate.
- 6. (Original) The formulation of claim 4 comprising between 1% and 20% w/w plasticizer.

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- 7. (Currently Amended) The formulation of claim 6 having wherein said coating has a viscosity of between 5 cP and 30 cP and a specific gravity of 1.054 g/ml plus or minus 0.050 g/ml.
- 8. (Original) The system formulation of claim 7 wherein the viscosity is between 5 cP and 20 cP.
 - 9. (Withdrawn) A formulation comprising (w/w):

20% to 35% ammonio methacrylate copolymer type B USP/NF;

1% to 10% triethyl citrate; and

60% to 70% water.

- 10. (Withdrawn) The formulation of claim 9 wherein the ammonio methacrylate copolymer is EUDRAGIT RS 30 D brand polymethylmethacrylate.
 - (Withdrawn) A formulation comprising (w/w) of: 11.

28% EUDRAGIT RS 30 D polymethylmethacrylate;

6% triethyl citrate; and

66% water.

(Withdrawn) A method for protecting pharmacologically-active substances applied 12. in a temporary coating to a surface of a tooth comprising:

applying a pharmacologically inert barrier coating of a polymethylmethacrylate and a plasticizer on top of the temporary coating containing the pharmacologically-active substance to serve as a temporary mechanical barrier against the washings of saliva and abrasion caused by eating food.

13. (Withdrawn) The method of claim 12 wherein the pharmacologically-active substance(s) comprise one or more active agents of the type known to reduce caries when applied to the tooth.

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14. (Withdrawn) The method of claim 13 wherein the pharmacologically-active substances are selected from the group consisting of chlorhexidine and fluoride.

- 15. (Withdrawn) The method of claim 12 wherein the polymethylmethacrylate is water-dispersed and the plasticizer is a pharmaceutical grade plasticizer selected from the group consisting of triethyl citrate, dibutyl sebacate, dibutyl phthalate, and diethyl phthalate.
- 16. (Withdrawn) The method of claim 12 wherein the polymethylmethacrylate is an ammonio methacrylate copolymer, type B USP/NF.
- 17. (Withdrawn) The method of claim 16 wherein the ammonio methacrylate copolymer is EUDRAGIT RS 30 D brand polymethylmethacrylate.
- 18. (Withdrawn) A method of preventing or reducing the incidence of caries in teeth, comprising the steps of:
 - a. applying a liquid coating of pharmacologically-active substances of the type used to reduce caries to a tooth surface; and
 - applying a pharmacologically inert barrier coating of a polymethylmethacrylate b. and a plasticizer on top of the coating containing the pharmacologically-active substance to serve as a temporary mechanical barrier against the washings of saliva and the abrasion from eating foods.